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**Company Location**

33 W. Thomas Rd, Suite 100  
Phoenix, AZ 85013

**Contact Numbers**

Tel: (602) 277-9256  
Fax: (602) 277-7150

**Contact:**

David Ratner

**Title:** Site Director

[davidratner@radiantresearch.com](mailto:davidratner@radiantresearch.com)

## **Overview**

Radiant Research, a Site Management Organization (SMO), conducts Phase I-IV human clinical trials for the biopharmaceutical industry through our nationwide network of clinical research facilities. Radiant Research employs nearly 1,000 research professionals throughout our 50+ wholly-owned clinical research facilities.

Radiant provides a critical service to biopharmaceutical companies by recruiting study participants into clinical trials, conducting the clinical trial protocol and collecting the data necessary to support the safety and efficacy of the drug company's product to the FDA. Our client list includes every major pharmaceutical company and contract research organization and many biotechnology and medical device companies. Radiant's clinical research sites have successfully completed more than 8,000 research studies in several areas of study.

## **Areas of Study**

Radiant's areas of study include: asthma and allergy, cardiology, central nervous system, dermatology, endocrinology and metabolism, gastroenterology, infectious disease, neurology, nutrition, over the counter/consumer products, pediatrics, preventative cardiology, psychiatry, pulmonology, rheumatology and musculoskeletal, urology, vaccines and women's health.

## **Our Mission**

It is our mission to dramatically improve the process of developing new drugs and medical devices. By assembling a group of the country's leading research physicians (investigators) in one unified organization, we are raising the standards of clinical studies. Human clinical trials represent two-thirds of the cost and over half of the time involved in bringing a drug to market. Radiant helps to reduce the cost and time of bringing new drugs to market by:

- **Initiating studies rapidly** through our wholly-owned, nationwide network of the most experienced, consistent and reliable clinical trial sites in the US.
- **Simplifying the administration** of clinical trials through the use of centralized institutional review boards, timely regulatory submissions, a single contract for multiple investigative sites and project management functions.
- **Accelerating the conduct** of clinical trials through our centralized participant recruitment services which professionally and cost-effectively match study participants with clinical trials.
- **Improving the reliability and accuracy** of clinical data through company-wide standard operating procedures and investigator and coordinator training and quality programs. Radiant Research is committed to bringing important new therapies to market more quickly, safely and efficiently.

## **Facility Description**

Radiant Research, Phoenix was founded in 1972 as Mary L. Wilson Clinical Research Center. Today the center occupies a 7,700 square foot medical facility entirely dedicated to clinical research.